



Atty. Dkt. No. 017371-0109

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

Applicant: Nicholas MANOLIOS  
Title: T CELL ANTIGEN RECEPTOR  
PEPTIDES  
Appl. No.: 09/202,305  
Filing Date: 3/22/1999  
Examiner: Anish Gupta  
Art Unit: 1654

**AMENDMENT AND REPLY UNDER 37 CFR 1.111**

Mail Stop Amendment  
Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

Sir:

This communication is responsive to the Non-Final Office Action dated May 5, 2004, concerning the above-referenced patent application.

Amendments to the Claims are reflected in the listing of claims which begins on page 2 of this document.

Remarks/Arguments begin on page 5 of this document.

Please amend the application as follows:

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**Amendments to the Claims**

This listing of claims will replace all prior versions and listings of claims in the application.

**Listing of Claims**

Claim 1. (Currently Amended) A peptide which inhibits T-cell antigen receptor (TCR) function, wherein the peptide is of the following formula:

R1-X-Z-X-R2 in which

X is a hydrophobic amino acid or a hydrophobic peptide sequence consisting of between 2 and 6 amino acids,

Z is a charged amino acid

R1 is NH<sub>2</sub> and

R2 is COOH,

wherein the hydrophobic peptide sequence does not include a charged amino acid, and

wherein the peptide is at least seven amino acids in length.

Claim 2. (Canceled)

Claim 3. (Previously Presented) The peptide according to claim 1 wherein at least 50% of the amino acids which make up the hydrophobic peptide sequence are hydrophobic amino acids.

Claim 4. (Previously Presented) The peptide according to claim 1 wherein Z is selected from Arg and Lys.

Claim 5. (Previously Presented). The peptide according to claim 1 which has the formula

NH<sub>2</sub>-Ile-Leu-Leu-Leu-Lys-Val-Ala-Gly-Phe-OH (SEQ ID NO. 6),

NH<sub>2</sub>-Ile-Leu-Leu-Leu-Lys-Val-Ala-Gly-OH (SEQ ID NO. 7),

NH<sub>2</sub>-Leu-Arg-Ile-Leu-Leu-Leu-Gly-Val-OH (SEQ ID NO. 8),

NH<sub>2</sub>-Leu-Gly-Ile-Leu-Leu-Leu-Lys-Val-OH (SEQ ID NO. 9),

NH<sub>2</sub>-Ile-Leu-Leu-Gly-Lys-Ala-Thr-Leu-Tyr-OH (SEQ ID NO. 10),  
NH<sub>2</sub>-Met-Gly-Leu-Arg-Ile-Leu-Leu-OH (SEQ ID NO. 11), or  
NH<sub>2</sub>-Leu-Leu-Met-Thr-Leu-Arg-Leu-Trp-Ser-Ser-COOH (SEQ ID NO. 12).

**Claim 6.** (Previously Presented) The peptide according to claim 1 wherein Z is selected from aspartic acid and glutamic acid.

**Claim 7.** (Original) A peptide according to claim 6 wherein the peptide has the formula

NH<sub>2</sub>-Ile-Ile-Val-Thr-Asp-Val-Ile-Ala-Thr-Leu-OH,  
NH<sub>2</sub>-Ile-Val-Ile-Val-Asp-Ile-Cys-Ile-Thr-OH, or  
NH<sub>2</sub>-Phe-Leu-Phe-Ala-Glu-Ile-Val-Ser-Ile-OH.

**Claim 8.** (Previously Presented) A peptide which inhibits TCR function, wherein the peptide is derived from the TCR- $\alpha$  intracellular chain and comprises the formula:

NH<sub>2</sub>-Ala-Gly-Phe-Asn-Leu-Leu-Met-Thr-COOH (SEQ ID NO. 16).

**Claim 9.** (Withdrawn) A peptide which inhibits TCR function, wherein the peptide is of the following formula:-

R<sub>1</sub>-A-B-C-R<sub>2</sub> in which

A is a peptide sequence of between 0 and 5 amino acids;

B is cysteine;

C is a peptide sequence of between 2 to 10 amino acids;

R<sub>1</sub> is NH<sub>2</sub>; and

R<sub>2</sub> is COOH.

**Claim 10.** (Withdrawn) A peptide according to claim 9 wherein A is a peptide sequence comprising 5 amino acids.

**Claim 11.** (Withdrawn) A peptide according to claim 9 wherein C is a peptide sequence of 4 or 5 amino acids and includes at least one hydrophobic amino acid.

**Claim 12.** (Previously Presented). A peptide which inhibits T-Cell antigen receptor function wherein the peptide has the formula:

NH<sub>2</sub>-Tyr-Gly-Arg-Ala-Asp-Cys-Gly-Ile-Thr-Ser-OH (SEQ ID NO. 17), or  
NH<sub>2</sub>-Trp-Gly-Arg-Ala-Asp-Cys-Gly-Ile-Thr-Ser-OH (SEQ ID NO. 18), or  
NH<sub>2</sub>-Tyr-Gly-Arg-Ala-Asp-Cys-Ile-Thr-Ser-OH (SEQ ID NO. 19), or  
NH<sub>2</sub>-Ser-Ser-Asp-Val-Pro-Cys-Asp-Ala-Thr-Leu-Thr-OH (SEQ ID NO. 20).

**Claim 13.** (Previously Presented) A therapeutic composition active against disorders in which T-cells are involved or recruited comprising a peptide as claimed in claim 1 and a pharmaceutically acceptable carrier.

**Claim 14.** (Withdrawn) A method of treating a subject suffering from a disorder in which T-cells are involved or recruited, the method including administering to the subject a therapeutically effective amount of the composition as claimed in claim 11.

**Claim 15.** (Canceled)

Please direct all correspondence to the undersigned attorney or agent at the address indicated below.

Respectfully submitted,

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